

**K050877 CORONARY SINUS DEFLECTABLE MAPPING
CATHETER, 4 FR, 5 FR, 6 FR & 7 FR**Jun 24, 2005
79 days to decisionK050877 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k050877/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Apr 6, 2005
Decision date	Jun 24, 2005
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biosense Webster, Inc.
Location	Irvine, CA, US
Contact	MICHAEL JOHNSON
Website	https://www.jnjmedtech.com
510(k) history	73 submissions · 73 cleared · 1999-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k050877/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026